

PACS1 gene

phosphofurin acidic cluster sorting protein 1

Normal Function

The *PACS1* gene provides instructions for making a protein called phosphofurin acidic cluster sorting protein 1 (PACS1). The PACS1 protein is found in a complex network of membranes known as the trans-Golgi network, which sorts proteins and other molecules and sends them to their intended destinations inside or outside the cell. Within the trans-Golgi network, this protein helps transport certain molecules and proteins. The PACS1 protein is most active during development before birth.

Health Conditions Related to Genetic Changes

PACS1 syndrome

At least two mutations in the *PACS1* gene have been found to cause *PACS1* syndrome. This condition is characterized by intellectual disability, speech and language problems, and a distinct facial appearance. Many affected individuals have additional neurological, behavioral, and health problems. The most common mutation, which occurs in nearly everyone with *PACS1* syndrome, results in the production of a protein with the protein building block (amino acid) arginine replaced with the amino acid tryptophan at position 203 (written as Arg203Trp or R203W).

PACS1 gene mutations are thought to impair the protein's ability to aid in the transport of molecules and proteins. Such an impairment likely results in the accumulation or misplacement of these substances within cells. The accumulated molecules and proteins may interfere with the function of the protein produced from the normal copy of the *PACS1* gene, further disrupting the placement of these substances.

Research suggests that impaired PACS1 protein function disrupts normal development of structures in the face, leading to a distinct facial appearance. It is likely that the development of other body systems are similarly affected by impaired PACS1 protein function, leading to other signs and symptoms of *PACS1* syndrome, but more research is needed to understand the mechanisms.

Other Names for This Gene

- cytosolic sorting protein PACS-1, human

- FLJ10209
- KIAA1175
- PACS-1
- phosphofurin acidic cluster sorting protein 1, human

Additional Information & Resources

Tests Listed in the Genetic Testing Registry

- Tests of PACS1 ([https://www.ncbi.nlm.nih.gov/gtr/all/tests/?term=55690\[geneid\]](https://www.ncbi.nlm.nih.gov/gtr/all/tests/?term=55690[geneid]))

Scientific Articles on PubMed

- PubMed (<https://pubmed.ncbi.nlm.nih.gov/?term=%28%28PACS1%5BTIAB%5D%29+OR+%28PACS-1%5BTIAB%5D%29%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+3600+days%22%5Bdp%5D%29%29%29>)

Catalog of Genes and Diseases from OMIM

- PHOSPHOFURIN ACIDIC CLUSTER SORTING PROTEIN 1; PACS1 (<https://omim.org/entry/607492>)

Gene and Variant Databases

- NCBI Gene (<https://www.ncbi.nlm.nih.gov/gene/55690>)
- ClinVar ([https://www.ncbi.nlm.nih.gov/clinvar?term=PACS1\[gene\]](https://www.ncbi.nlm.nih.gov/clinvar?term=PACS1[gene]))

References

- Lusk L, Smith S, Martin C, Taylor C, Chung W. PACS1 Neurodevelopmental Disorder. 2020 Jul 16. In: Adam MP, Feldman J, Mirzaa GM, Pagon RA, Wallace SE, Bean LJH, Gripp KW, Amemiya A, editors. GeneReviews(R) [Internet]. Seattle (WA): University of Washington, Seattle; 1993-2024. Available from <http://www.ncbi.nlm.nih.gov/books/NBK559434/> Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/32672908>)
- Schuurs-Hoeijmakers JH, Landsverk ML, Foulds N, Kukolich MK, Gavrilova RH, Greville-Heygate S, Hanson-Kahn A, Bernstein JA, Glass J, Chitayat D, Burrow TA, Husami A, Collins K, Wusik K, van der Aa N, Kooy F, Brown KT, Gadzicki D, Kini U, Alvarez S, Fernandez-Jaen A, McGehee F, Selby K, Tarailo-Graovac M, Van Allen M, van Karnebeek CD, Stavropoulos DJ, Marshall CR, Merico D, Gregor A, Zweier C, Hopkin RJ, Chu YW, Chung BH, de Vries BB, Devriendt K, Hurles ME, Brunner HG;

DDDstudy. Clinical delineation of the PACS1-related syndrome--Report on 19 patients. Am J Med Genet A. 2016 Mar;170(3):670-5. doi: 10.1002/ajmg.a.37476. Epub 2016 Feb 3. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/26842493>)

- Schuurs-Hoeijmakers JH, Oh EC, Vissers LE, Swinkels ME, Gilissen C, Willemsen MA, Holvoet M, Steehouwer M, Veltman JA, de Vries BB, van Bokhoven H, de Brouwer AP, Katsanis N, Devriendt K, Brunner HG. Recurrent de novo mutations in PACS1 cause defective cranial-neural-crest migration and define a recognizable intellectual-disability syndrome. Am J Hum Genet. 2012 Dec 7;91(6):1122-7. doi:10.1016/j.ajhg.2012.10.013. Epub 2012 Nov 15. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/23159249>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3516611/>)
- Stern D, Cho MT, Chikarmane R, Willaert R, Retterer K, Kendall F, Deardorff M, Hopkins S, Bedoukian E, Slavotinek A, Schrier Vergano S, Spangler B, McDonald M, McConkie-Rosell A, Burton BK, Kim KH, Oundjian N, Kronn D, Chandy N, Baskin B, Guillen Sacoto MJ, Wentzensen IM, McLaughlin HM, McKnight D, Chung WK. Association of the missense variant p.Arg203Trp in PACS1 as a cause of intellectual disability and seizures. Clin Genet. 2017 Aug;92(2):221-223. doi:10.1111/cge.12956. Epub 2017 Jan 23. Citation on PubMed (<https://pubmed.ncbi.nlm.nih.gov/28111752>) or Free article on PubMed Central (<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5513756/>)

Genomic Location

The *PACS1* gene is found on chromosome 11 (<https://medlineplus.gov/genetics/chromosome/11/>).

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